



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61B 17/34, A61M 29/00	A1	(11) International Publication Number: WO 00/10471 (43) International Publication Date: 2 March 2000 (02.03.00)
(21) International Application Number: PCT/US99/19073 (22) International Filing Date: 19 August 1999 (19.08.99) (30) Priority Data: 60/097,126 19 August 1998 (19.08.98) US 60/106,733 2 November 1998 (02.11.98) US Not furnished 18 August 1999 (18.08.99) US (71) Applicant: GENESIS MEDICAL TECHNOLOGIES, INC. [US/US]; P.O. Box 246, Redwood City, CA 94064 (US). (72) Inventors: FULTON, Richard, Eustis, III; 1556 Wellington Avenue, Grand Junction, CO 81501 (US). DUBRUL, William, R.; Suite A-7, 1 Uccelli Boulevard, Redwood City, CA 94063 (US). (74) Agents: HANN, James, F. et al.; Townsend and Townsend and Crew LLP, 8th Floor, Two Embarcadero Center, San Francisco, CA 94111-3834 (US).		(81) Designated States: JP, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: TARGET TISSUE LOCALIZATION DEVICE AND METHOD <div style="text-align: center;"> </div> (57) Abstract <p>A target tissue localization device (28) has outer and inner members (30, 32) which are longitudinally movable relative to one another. An anchor (42) is secured to the outer and inner members and is movable from a radially-contracted configuration to a radially-expanded configuration. The anchor may be a tubular mesh anchor. The anchor may have a sufficiently large cross-sectional area when in the radially-expanded configuration to substantially eliminate inadvertent dislodgement of the anchor once radially-expanded. In use, the location of the target tissue (56) is determined and the distal end of the localization device is passed through the patient's skin (58) and to the target tissue to create a tissue track (60) between the patient's skin and the target tissue. The target tissue includes near and far sides (62, 64). The anchor element, located at the target tissue is expanded to an expanded configuration. The method the anchor element may be located on the far side of the target tissue. The localization device may be selected with an anchor which is palpable for relocation of the target tissue.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand		
CM	Cameroon		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

TARGET TISSUE LOCALIZATION DEVICE AND METHOD

BACKGROUND OF THE INVENTION

In the U.S. alone approximately one million women will have breast
5 biopsies because of irregular mammograms and palpable abnormalities. Biopsies can be
done in a number of different ways for non-palpable lesions, including surgical excisional
biopsies and stereotactic and ultrasound guided needle breast biopsies. In the case of
image directed biopsy, the radiologist or other physician takes a small sample of the
10 irregular tissue for laboratory analysis. If the biopsy proves to be malignant, additional
surgery (typically a lumpectomy or a mastectomy) is required. In the case of needle
biopsies, the patient then returns to the radiologist a day or two later where the biopsy site
(the site of the lesion) is relocated by method called needle localization, a preoperative
localization in preparation for the surgery.

Locating the previously biopsied area after surgical excision type of biopsy
15 is usually not a problem because of the deformity caused by the surgery. However, if the
biopsy had been done with an image directed needle technique, as is common, help in
relocating the biopsy site is needed. One procedure to permit the biopsy site to be
relocated by the radiologist during preoperative localization is to leave some of the
suspicious calcifications; this has its drawbacks.

20 Another way to help the radiologist relocate the biopsy site involves the
use of a small metallic surgical clip, such as those made by Biopsys. The metallic clip
can be deployed through the biopsy needle, and is left at the biopsy site at the time of the
original biopsy. With the metallic clip as a guide, the radiologist typically inserts a barbed
or hooked wire, such as the Hawkins, Kopans, Homer, Sadowski, and other needles, back
25 into the patient's breast and positions the tip of the wire at the biopsy site using
mammography to document the placement. The patient is then taken to the operating
room with the needle apparatus sticking out of the patient's breast. While the clip
provides a good indication of the biopsy site to the radiologist during preoperative
localization, the clip remains permanently within the 80% of patients with benign
30 diagnoses. Also, because the clip is necessarily attached to a single position at the
periphery of the biopsy site, rather than the center of the biopsy site, its location may
provide a misleading indication of the location of diseased tissue during any subsequent
medical intervention. The clip is also relatively expensive. In addition, the soft nature of

breast tissue permits the tip of the barbed or hooked needle to be relatively easily dislodged from the biopsy site. The surgeon typically pulls on the needle to help locate the area to be removed. This pulling motion during the excision may be responsible more than any other single factor for the movement of the needle. Additionally, these devices
5 are not easily felt by external palpation and the tip is difficult to locate without dissection into the tissues; this prevents a surgical approach which may be more cosmetically advantageous and surgically appropriate than dissecting along the tract of the needle.

Another localization method involves the use of laser light from the tip of a optical fiber connected to a laser. A pair of hooks at the tip of the optical fiber secures
10 the tip at the biopsy site; the glow indicates the position of the tip through several centimeters of breast tissue. This procedure suffers from some of the same problems associated with the use of barbed or hooked wires. Another preoperative localization procedure injects medical-grade powdered carbon suspension from the lesion to the skin surface. This procedure also has certain problems, including the creation of
15 discontinuities along the carbon trail.

SUMMARY OF THE INVENTION

The present invention is directed to a target tissue localization device including an elongate placement device having first and second members being
20 longitudinally movable relative to one another. An anchor is secured to the first and second members and is movable from a first, radially contracted configuration to a second, radially expanded configuration. The placement device and anchor are sized and configured for use in localization of target tissue. According to one aspect of the invention the anchor is a tubular mesh anchor. According to another aspect of the
25 invention the anchor has a sufficiently large cross-sectional area when in the second, radially expanded configuration to substantially eliminate inadvertent dislodgment of the anchor once radially-expanded.

A further aspect of the invention relates to a method for localizing target tissue including determining the location of the target tissue and passing the distal end of
30 an elongate localization device through the patient's skin and to the target tissue to create a tissue track between the patient's skin and the target tissue. The target tissue includes a near side toward the patient's skin and a far side opposite the near side. The localization device includes an anchor element secured to a placement device, the placement device extending from the anchor element along the tissue track and through the patient's skin.

The anchor element is located at the target tissue and expanded to an expanded configuration. According to one aspect of the method the anchor element is located on the far side of the target tissue. According to another aspect of the method, the localization device is selected with an anchor which is palpable at the patient's target
5 tissue so that the target tissue can be relocated at least in part by palpation of the patient to locate the anchor; according to this aspect of the invention, at least a sample of the relocated target tissue is removed. The elongate localization device can be used as a guide or guide wire for further diagnostic of therapeutic procedures.

In preferred embodiment of the invention the anchor is a tubular mesh
10 anchor which can be placed into a disk-like shape when in the fully expanded configuration or in a more flattened spherical shape, that is not as flat as the disk-like shape, which may aid in palpability. The anchor may also be in other forms such as a Malecot-type of radially expandable anchor. Another anchor embodiment could be deployed from the end of a hollow sheath and include numerous resilient curved
15 projections which extend generally radially outwardly to fully surround the sheath.

The invention may find particular utility as an aid in the removal of small, difficult-to-grasp body structures, such as lymph nodes or other tissue that is difficult to grasp. The anchor could be positioned behind or inside a lymph node targeted for removal; lymph node removal can be aided by leaving the anchor in its expanded position
20 so that after the surgeon cuts down to the lymph node, the lymph node, some surrounding tissue and the entire localization device can be removed for further analysis.

In some situations the anchor may be left in a partially or fully radially-expanded condition and used to pull the target tissue from the patient, typically with the aid of the surgeon opening the tissue track. For example, for tumor removal of the breast,
25 the anchor may be used in a partially or fully expanded condition so that the tumor and surrounding margins can be removed during lumpectomy. Similar procedures can be used in regions other than the breast.

The invention may also include structure which may enhance ultrasound visibility. Invention may include certain treatment, grooves, slits, or other structure to
30 enhance the ultrasonic visibility of the placement device or the delivery needle, or both. Also, the mesh anchor may provide sufficient reflectivity so that treatment of the placement device and delivery needle is not required. Other techniques for enhancing remote visualization, such as radiopaque markers, can also be used.

The invention is primarily related to mammographic localization, but may find utility with other areas of imaging including ultrasound, thoracoscopy, nuclear medicine, MRI, computed tomography, plain film and any other image guided technology.

5 The present invention provides a simple, relatively inexpensive, very stable localization device to locate a lesion or other target tissue, typically within the breast of a patient, for subsequent biopsy, excision, intervention, or other purpose. The localization device is easy to insert, deploy, reposition if needed, and remains anchored at the target site substantially better than conventional localization needles. Further, the
10 invention aids relocation of the target tissue by being configured to permit location by palpation by the surgeon in the operating suite. This permits the surgeon to proceed using much less radical techniques, which may be both cosmetically and surgically advantageous.

 Other features and advantages of the invention will appear from the
15 following description in which the preferred the embodiments have been set forth in detail in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

 Fig. 1 illustrates a conventional hollow insertion needle and hook wire;
20 Fig. 2 illustrates the hook wire of Fig. 1 inserted into the insertion needle with the barb extending through the opening in the insertion needle;
 Fig. 3 is an overall view of a localization assembly made according to the invention with the anchor in a radially expanded, deployed condition;
 Fig. 4 is a partial cross-sectional view of the distal portion of the
25 localization device of Fig. 2 with the anchor in a radially contracted configuration;
 Fig. 5 illustrates the localization device of Fig. 4 with the anchor moved to a radially expanded configuration by the movement of the distal ends of the outer sleeve member and the inner wire member towards one another;
 Fig. 5A is a cross-sectional view taken along line 5A-5A of Fig. 5;
30 Fig. 6 illustrates a further embodiment of the localization device of Fig. 4 in which the inner wire member has been replaced by an inner shaft member having a pointed distal end to potentially eliminate the need for the delivery needle of Fig. 3;
 Fig. 7 illustrates the device of Fig. 6 with the anchor in a first radially expanded condition, the anchor forming a somewhat flattened spherical shape;

Fig. 8 illustrates the localization device of Fig. 7 with the anchor in a fully radially expanded configuration in the shape of a generally flat disk, the anchor illustrated positioned on the far side of a lesion or other target tissue;

Fig. 9 illustrates an alternative embodiment of the localization assembly of Fig. 3 in which the localization device of Fig. 3 has been replaced by a localization device having an inner shaft member and an outer shaft member, the outer shaft member having a series of longitudinally extending slits formed at its distal end;

Fig. 10 illustrates the localization device of Fig. 9 with the inner shaft member pulled relative to the outer sleeve member causing the sleeve segments to expand radially to form a Malecot-type anchor; and

Fig. 11 illustrates a further alternative embodiment of the invention in which the anchor is in the form of an expandable balloon, the balloon mounted to the distal end of a shaft, the balloon-type anchor and distal shaft end housed within a delivery needle;

Fig. 11A illustrates the embodiment of Fig. 11 which the balloon-type anchor is in its radially-contracted condition and is positioned past the sharpened tips of the delivery needle; and

Fig. 11B illustrates the shaft and balloon-type anchor of Fig. 11A with the balloon-type anchor in its radially-expanded, deployed condition.

DESCRIPTION OF THE SPECIFIC EMBODIMENTS

Fig. 1 illustrates a conventional hollow insertion needle 2 placed next to its associated hook wire 4. Insertion needle 2 has a sharpened tip 6 and an opening 8 set back away from tip 6. Hook wire 4 includes a shaft 10 and a barb 12 extending off the distal end of the shaft. Insertion needle 2 is relatively stiff or rigid while hook wire 4 is flexible but resilient. Fig. 2 illustrate the localization assembly 14 made up of the insertion needle 2 and hook wire 4 of Fig. 1 with the barb 12 extending from opening 8. The condition of localization assembly 14 in Fig. 2 is that which would be after tip 6 of insertion needle 2 has been properly placed relative to the target tissue and hook wire 4 has been pulled proximally causing barb 12 to extend from opening 8 and pass into the patient's tissue. As discussed above, this reliance on a thin barb 12 to ensure that assembly 14 does not move after placement is not particularly well-founded; breast tissue is generally quite soft and permits barb 12 to be easily dislodged from its initial position thus providing the surgeon a false indication of the target tissue location.

Fig. 3 illustrates a localization assembly 20 made according to the invention. Assembly 20 includes broadly a hollow delivery needle 22 having a sharpened tip 24 at its distal end and a hub 26 at its proximal end. Needle 22 houses a localization device 28, see Figs. 4, 5 and 5A, the localization device including an outer sleeve member 30 and an inner wire member 32. The distal ends 34, 36 of outer sleeve member 30 and inner wire member 32 are secured to the proximal and distal ends 38, 40 of a tubular mesh anchor 42, respectively. The relative positions of hollow delivery needle 22 and outer sleeve member 30 can be locked in place by a needle/sleeve locking device 44 while the relative positions between outer sleeve member 30 and inner wire member 32 can be secured by a sleeve/wire locking device 46. Anchor 42 is moved from its radially contracted condition of Fig. 4 to its radially expanded condition of Figs. 3 and 5 by pulling on inner wire member 32 while maintaining outer sleeve member 30 in place, or by pushing on outer sleeve member 30 while maintaining inner wire member 32 in place, or by both pulling on inner wire 32 and pushing on outer sleeve 30 as suggested by the arrows in Fig. 5.

Localization assembly 20 is preferably made from conventional bio-compatible materials, such as stainless steel for delivery needle 22, outer sleeve member 30 and inner wire member 32. Anchor 42 is preferably made of stainless steel or NiTi or a polymer such as Kynar, polyester, nylon, etc., and may be coated with a material such as PTFE or other lubricious coating, such as hydrophilic or hydrophobic coating commonly used in medical devices, to help anchor 42 pass through the patient's tissue both when in the radially contracted condition of Fig. 4 and when moved to the radially expanded condition of Fig. 5. The mesh material of mesh anchor 42 preferably has a tight enough weave so that it provides substantial hindrance to passage of the patient's tissue, typically at a breast tissue, through the anchor when in the radially expanded condition of Figs. 3 and 5. Tubular mesh anchor 42 may be somewhat porous or it could be substantially impervious to fluid flow. Anchor 42 could be, for example, covered with a flexible elastic film to provide fluid impermeability in the expanded condition.

In one embodiment anchor 42 is made of wires, or filaments or yarns having a diameter of .13mm (.005 inch) and spacing of .025mm (.001 inch) in its contracted condition of Fig. 4. Mesh anchor 42 is preferably made from wires or other elements have diameters of about .025 to .38mm (.001 to .015 inch) and a contracted condition spacing of about .0025 to .051mm (.0001 to .002 inch). The length of anchor 42 is a large factor in determining how large anchor 42 is when in the radially expanded

condition of Figs. 5 and 5A. That is, a tubular anchor 42 having a relatively short length when in the radially contracted condition of Fig. 4 will have a smaller diameter D when in the radially expanded condition of Figs. 5 and 5A as opposed to an anchor with a longer axial length when in the radially contracted condition of Fig. 4.

5 The cross-sectional area D diameter of anchor 42 in the radially expanded condition of Figs. 5 and 5A, equal to $\pi D^2/4$, is preferably at least four times and more preferably at least eight times the cross-sectional area of anchor 42 in the radially contracted condition of Fig. 4, equal to $\pi d^2/4$. Another way of defining the enhanced stabilization created by anchor 42 is based upon the force it would take to substantially
10 dislodge the anchor from a deployed position within the patient's breast. Conventional hooked wire embodiments such as that of Figs. 1 and 2, which commonly become substantially dislodged when a longitudinally directed pulling force of only about .5 to 2 kg (1 to 4 pounds) is exerted on localization assembly 14. Anchor 42, due to its large surface area opposing any axially-directed force, requires a longitudinally directed pulling
15 force on outer sleeve 30 and inner wire member 32 to be substantially more than the force required to substantially dislodge a hooked wire localization needle to be substantially dislodged from breast tissue.

 An alternative embodiment of the localization assembly is shown in Fig. 6 with like reference numerals referring to like elements. Localization device 28A is
20 designed to potentially eliminate the need for the hollow delivery needle 22 of Fig. 3. Outer sleeve 30A is essentially the same with an inner shaft 32A replacing inner wire 32. The distal end 36A of inner shaft 32A is a sharpened point to aid piercing the patient's tissue. The tubular mesh anchor 42A is secured at its distal and proximal ends 40A and 38A to distal end 36A of inner shaft 32A and distal end 34A of outer sleeve 30A. Fig. 7
25 illustrates result of collapsing anchor 42A about half way resulting in a generally flatten spherical shape or ellipsoid of revolution shape. This shape does not create as large a cross-sectional area but may be useful in aiding palpability of anchor 42A. Fig. 8 illustrates the result of continuing to collapse inner mesh anchor 42A by pulling on inner shaft member 32A; doing so results in anchor 42A being similar in shape to anchor 42 of
30 Figs. 5 and 5A.

 A further embodiment of the invention is illustrated in Figs. 9 and 10. In Fig. 9, a localization assembly 20B is seen to include a delivery needle 22B housing a localization device 28B. Localization device 28B includes an outer sleeve member 30B

and an inner shaft member 32B. The distal end 34B of outer shaft member 30B has a member of axially-extending slits 48 formed therein to define a plurality, preferably six to ten, sleeve segments 50 therebetween. Pulling inner shaft member 32B relative to outer shaft member 30B causes sleeve segments 50 to buckle outwardly to create a Malecot-type anchor 52. To improve the effective cross-sectional area of anchor 52 in the radially-expanded condition, sleeve segments 50 could be surrounded by an expandable mesh or a film which would effectively cover or fill the regions between sleeve segments 50 when in the expanded condition of Figs. 10 and 10A.

The invention may be used in a manner similar to conventional localization devices. The location of the target tissue 56, see Fig. 8, is determined using, for example, mammography, ultrasound, or other conventional or unconventional techniques. Using the embodiment of Figs. 6-8 as an example, localization device 28A is passed through skin 58 to create a tissue path 60 between target tissue 56 and the skin. Target tissue 56 has a near side 62 towards skin 58 and a far side 64 away from near side 62, that is facing anchor 42A in Fig. 8. Once in the expanded condition of Fig. 8, localization device 28A is very resistant to being moved or dislodged so that anchor element 42A remains where positioned, in this example on the far side of target tissue 56. Locking device 46 is used to maintain the relative axial positions of outer sleeve member 30 and inner shaft member 32A. In example Fig. 8, localization device 28A is shown having pierced through target tissue 56. In appropriate cases localization device 28A could be positioned to one side of target tissue 56 with anchor 42A opposite far side 64. Distal end 36A could also be positioned opposite near side 62 so that the localization device 28A does not pierce target tissue 56 and anchor 42A lies opposite near side 62 rather than far side 64. The surgeon can relocate target tissue 56 using localization 28A in several ways. First, when anchor 42A is constructed for palpability, the surgeon may determine the location of target tissue 56 by palpation by sensing the presence and location of anchor 42A, anchor 42A being harder than the surrounding tissue. Remote visualization techniques could also be used to determine the location of anchor 42A and thus of target tissue 56. Appropriate therapeutic or diagnostic procedures, such as taking biopsy samples of target tissue 56 or completely removing target tissue 56, can then be accomplished. In some situations localization device 28A will remain in position until anchor 42A is transformed from its radially-expanded condition of Fig. 8 to its radially-contracted condition of Fig. 6, at which time device 28A can be removed from the patient. At other times anchor 42A will remain in its fully radially deployed condition as

localization device 28A is used to help pull target tissue 56 along tissue path 60, which may or may not occur after surgical intervention.

A further embodiment of the invention is shown in Figs. 11-11B. This embodiment uses a balloon-type anchor 70 at the distal end of a hollow delivery shaft 72. Anchor 70 is delivered to a target site while housed within a hollow delivery needle 22 as shown in Fig. 11. Once at the proper location, anchor 70 is exposed, as shown in Fig. 11A, by pushing on shaft 72, pulling on needle 22, or both. Fig. 11B shows balloon-type anchor 70 in its radially-expanded, deployed condition, a flattened spherical shape in this embodiment. Balloon anchor 70 may be made of elastic or inelastic material and can be made to assume a variety of shapes when in the expanded condition. Such shapes include the disk-like shapes Figs. 3, 5, 5A, the flattened spherical shaped Figs. 7 and 11B, star-shaped with radially extending fingers (not shown) or other shapes suitable for the surrounding tissue in which it is deployed and the circumstances of its use.

Modification and variation can be made to the disclosed embodiments without departing from the subject of the invention as defined in the following claims.

Any and all patents, patent applications and printed publications referred to above are incorporated by reference.

WHAT IS CLAIMED IS:

- 1 1. A target tissue localization device comprising:
2 an elongate placement device comprising:
3 a first member having a distal portion; and
4 a second member carried by the first member and being
5 longitudinally movable relative thereto;
6 an anchor having first and second portions secured to the first and second
7 members, respectively;
8 said anchor being a tubular mesh anchor movable from a first, radially
9 contracted configuration to a second, radially expanded configuration;
10 said placement device and anchor being sized and configured for use in
11 localization of target tissue.
- 1 2. The device according to claim 1 wherein the first member is a
2 tubular member and the second member is slidably housed within the first member.
- 1 3. The device according to claim 2 wherein the second member
2 comprises a flexible wire and the first member comprises a stiff tube.
- 1 4. The device according to claim 2 wherein the first and second
2 members are stiff members.
- 1 5. The device according to claim 1 wherein the first and second
2 portions of the anchor are proximal and distal portions of the anchor.
- 1 6. The device according to claim 5 wherein the second member has a
2 pointed distal end.
- 1 7. The device according to claim 5 wherein the first and second
2 members have distal ends and the first and second portions of the anchor are secured to
3 the distal ends of the first and second members, respectively.
- 1 8. The device according to claim 5 wherein the first and second
2 members have distal ends and the first and second portions of the anchor are secured to
3 the distal ends of the first and second members, respectively.

1 9. The device according to claim 1 wherein:
2 said anchor has first and second cross-sectional areas, lying in radially-
3 oriented planes, when in the first and second configurations, said second area being at
4 least four times said first area.

1 10. A target tissue localization device comprising:
2 an elongate placement device comprising:
3 a first member having a distal portion; and
4 a second member carried by the carrier member and being
5 longitudinally movable relative thereto;
6 an anchor having first and second portions secured to the first and second
7 members, respectively;
8 said anchor movable from a first, radially contracted configuration to a
9 second, radially expanded configuration;
10 said placement device and anchor being sized and configured for use in
11 localization of target tissue; and
12 said anchor having first and second cross-sectional areas, lying in radially-
13 oriented planes, when in the first and second configurations, said second area being at
14 least four times said first area.

1 11. A method for localizing a target tissue comprising:
2 determining the location of the target tissue;
3 passing the distal end of an elongate localization device through the
4 patient's skin and to the target tissue to create a tissue track between the patient's skin and
5 the target tissue, the target tissue comprising a near side towards the patient's skin and a
6 far side opposite the near side, the localization device comprising an anchor element
7 secured to a placement device, the placement device extending from the anchor element,
8 along the tissue track and through the patient's skin;
9 locating the anchor element on the far side of the target tissue; and
10 expanding the anchor element to an expanded configuration on the far side
11 of the target tissue.

1 12. The method according to claim 11 further comprising:
2 relocating the target tissue using the localization device; and

3 removing at least a sample of the relocated target tissue.

1 13. The method according to claim 11 further comprising selecting the
2 localization device having an anchor which is palpable at the patient's target tissue.

1 14. The method according to claim 13 wherein the relocating step is
2 carried out at least in part by palpation of the patient.

1 15. The method according to claim 13 wherein the selecting step
2 comprises selecting an anchor having first and second cross-sectional areas, lying in
3 radially-oriented planes, when in the first and second configurations, said second area
4 being at least four times said first area.

1 16. The method according to claim 11 wherein the location
2 determining step is carried out by at least one of x-ray, ultrasound, fluoroscope, MRI,
3 nuclear medicine, computer tomography, plain film and other image guided techniques.

1 17. The method according to claim 21 wherein the passing step is
2 carried out with the aid of a hollow introducer needle having a sharpened tip and housing
3 the placement device.

1 18. A method for localizing target tissue comprising:
2 determining the location of the target tissue;
3 selecting an elongate localization device having an anchor which is
4 palpable at the patient's target tissue when in an expanded configuration at the target
5 tissue, the localization device comprising a guide element extending from the anchor;
6 passing a distal end of the localization device through the patient's skin
7 and to the target tissue to create a tissue track between the patient's skin and the target
8 tissue, the guide element extending along the tissue track and through the patient's skin;
9 locating the anchor element at the target tissue;
10 expanding the anchor element from a contracted configuration to the
11 expanded configuration;
12 relocating the target tissue at least in part by palpation of the patient to
13 locate the anchor; and
14 removing at least a sample of the relocated target tissue.

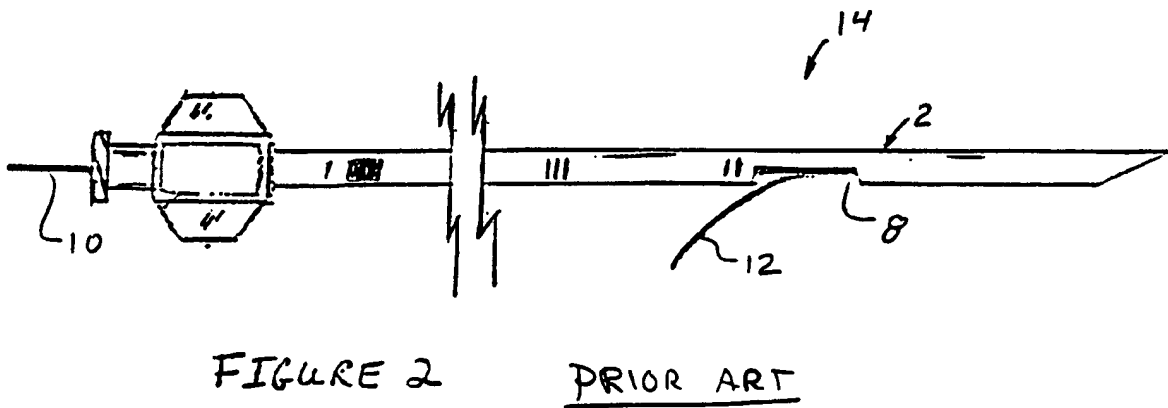
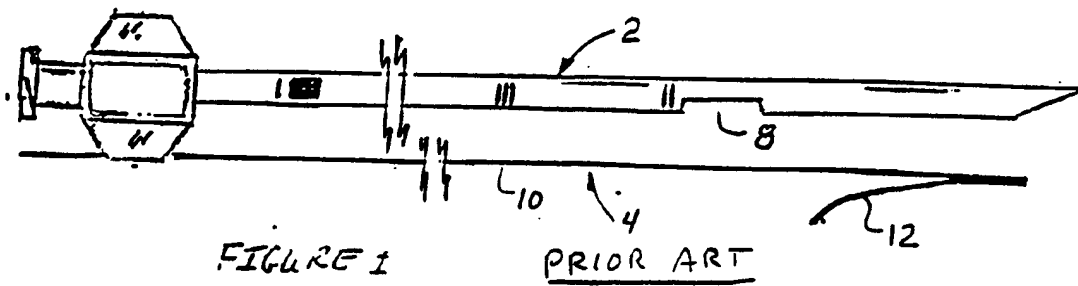
1 19. The method according to claim 18 wherein the expanding step
2 causes the anchor to assume a generally flattened spherical shape.

1 20. The method according to claim 18 wherein the selecting step
2 selects a Malecot-type anchor having a plurality of generally radially-extending segments
3 when in the expanded configuration.

1 21. The method according to claim 18 wherein the selecting step
2 selects a tubular mesh anchor.

1 22. The method according to claim 18 wherein the expanding step is
2 carried out so the anchor, when in the expanded configuration, has a cross-sectional area
3 at least four times the cross-sectional area of the anchor when in the contracted
4 configuration.

1 23. The method according to claim 18 wherein the selecting step
2 selects a balloon-type anchor.



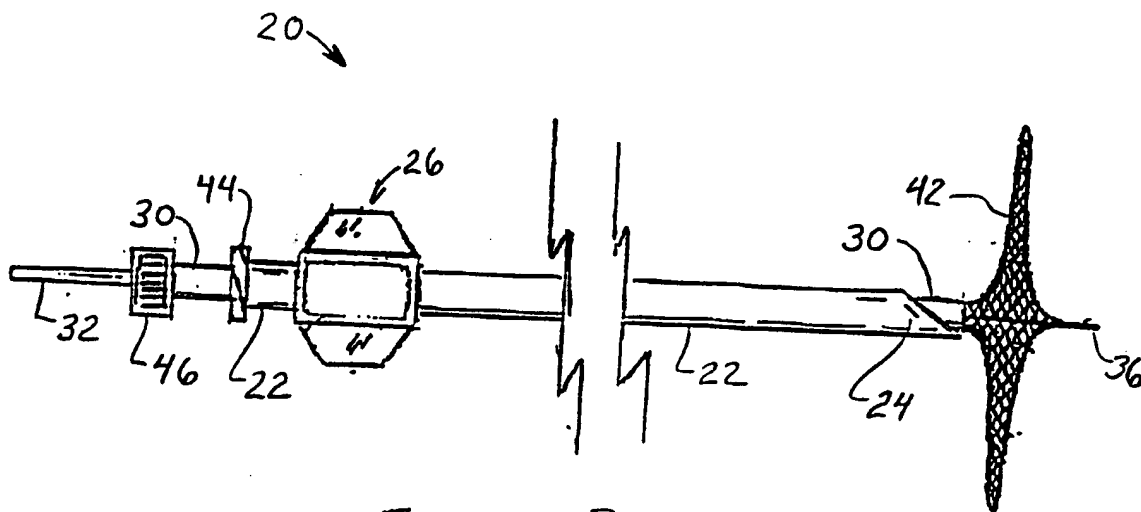


FIGURE 3

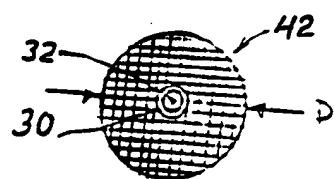
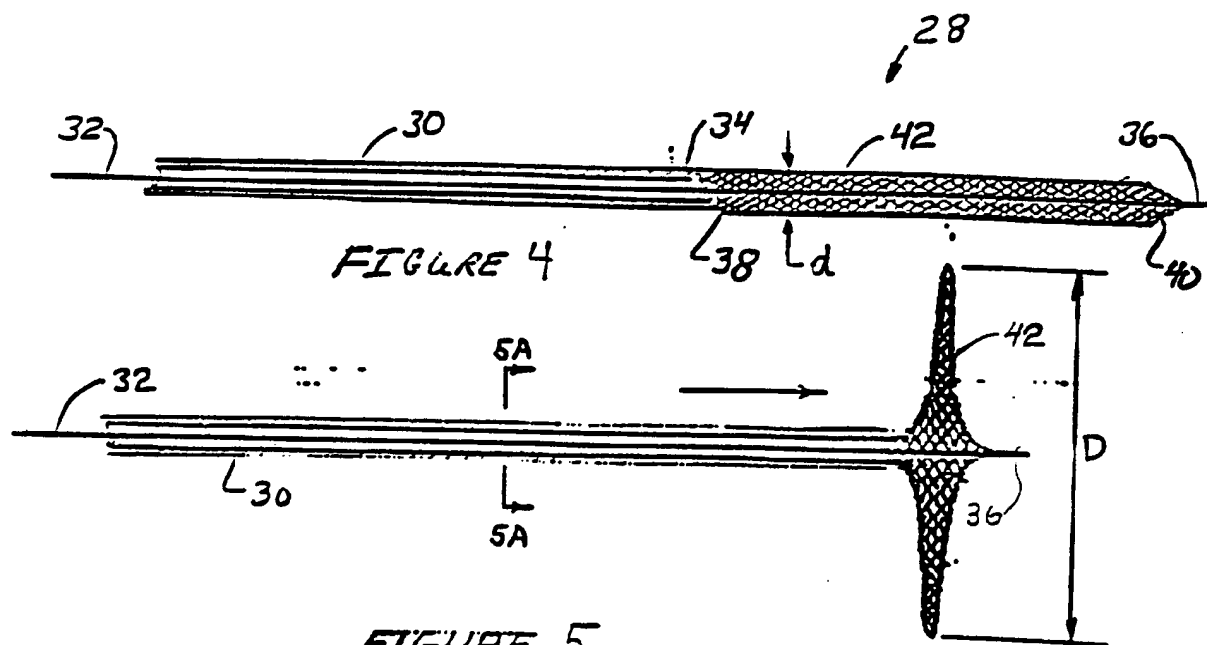
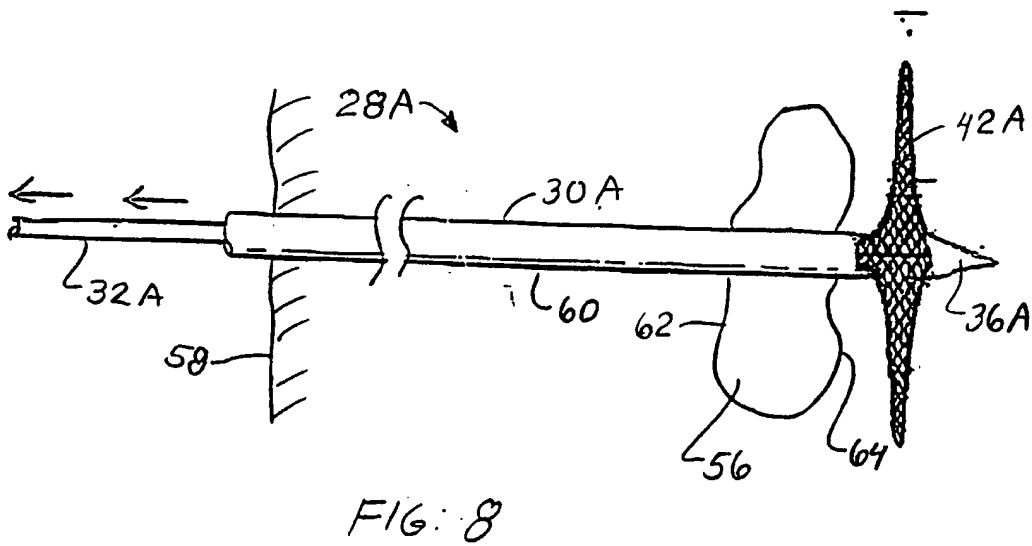
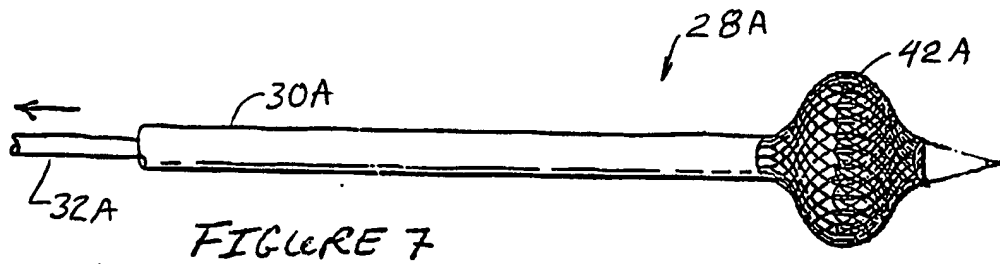
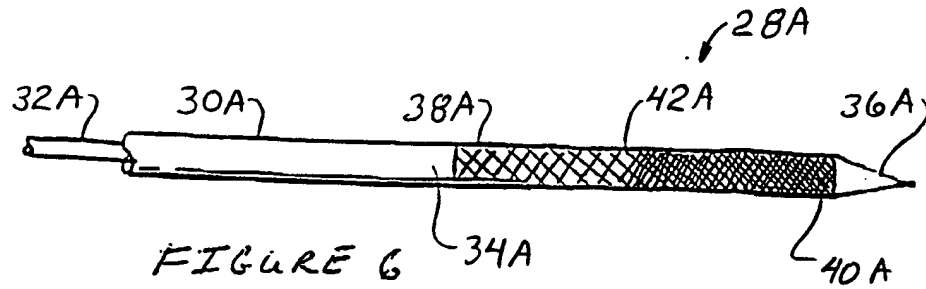
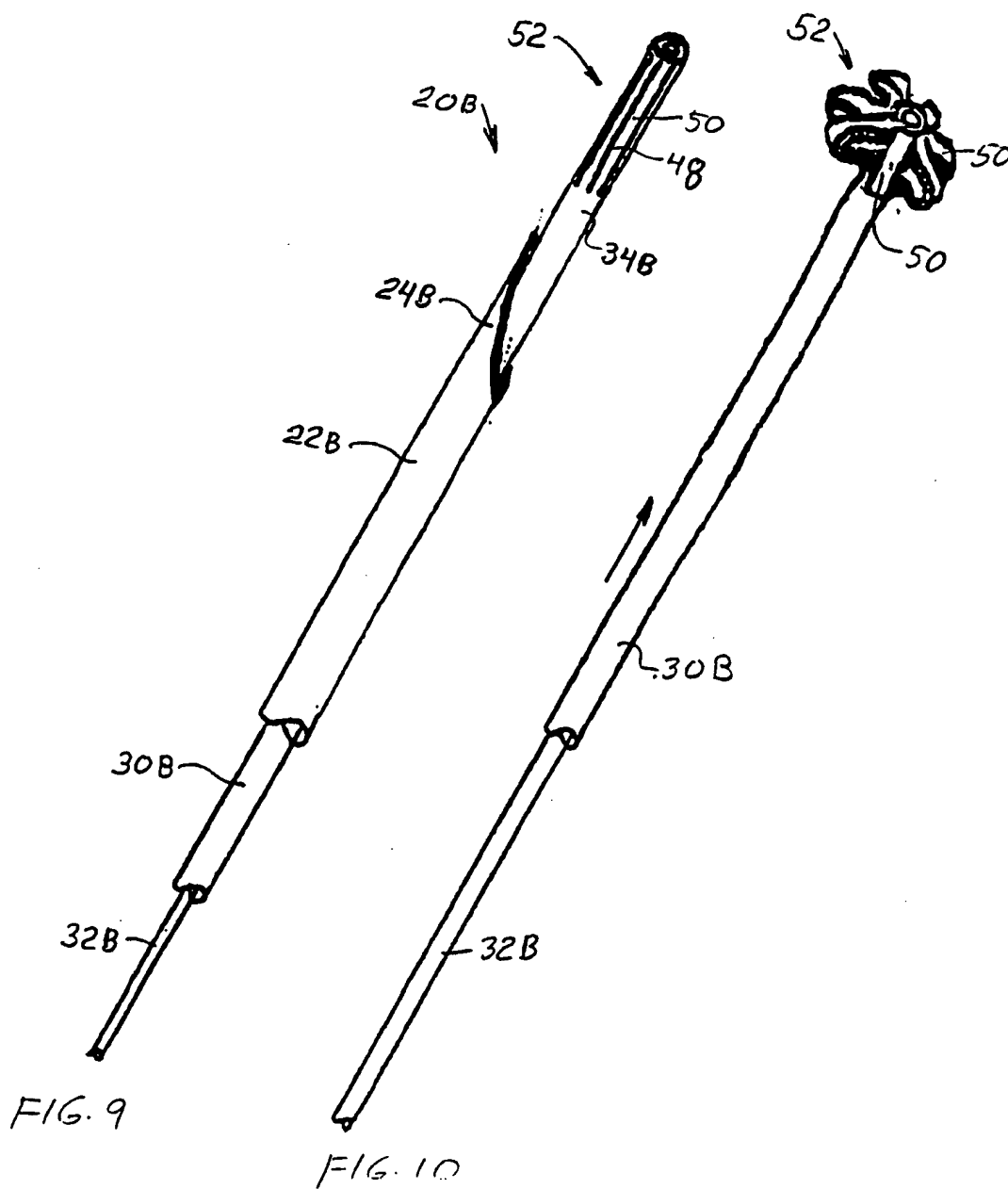


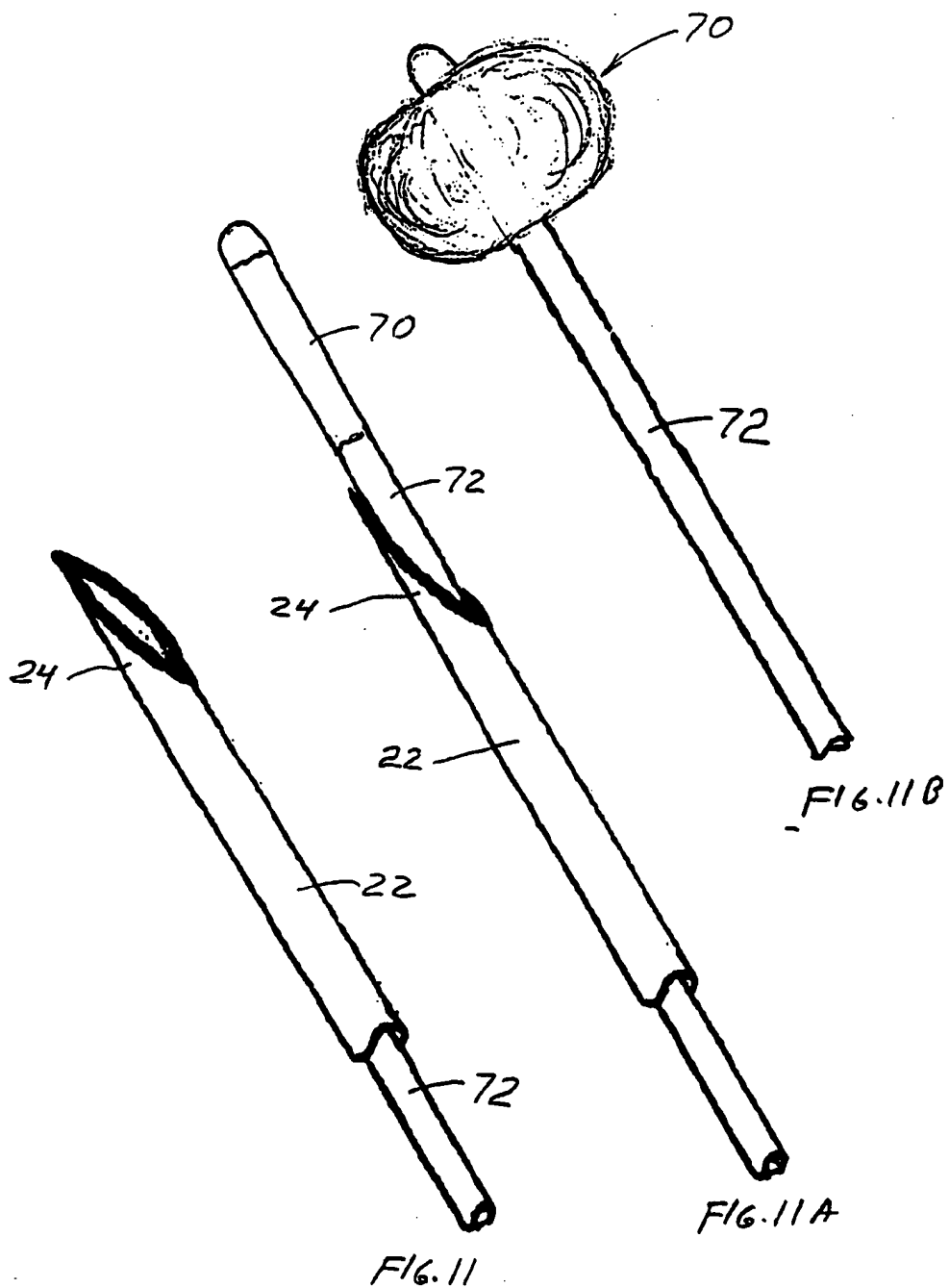
FIG. 5A

4/6





6/6



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US99/19073

A. CLASSIFICATION OF SUBJECT MATTER IPC(6) : A61B 17/34; A61M 29/00 US CL : 606/185, 192, 194, 197, 198 According to International Patent Classification (IPC) or to both national classification and IPC																				
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) U.S. : 606/185, 192, 194, 197, 198 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) APS Search Terms: needle, biopsy, anchor, breast.																				
C. DOCUMENTS CONSIDERED TO BE RELEVANT																				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.																		
X — Y	US 5,795,308 A (RUSSIN) 18 August 1998, col. 1, lines 11-14; col. 6, lines 10-16; and Figs. 1-25C.	11, 12, 16, 18, 20 13, 14, 17, 19, 21-23																		
Y	US 5,183,463 A (DEBBAS) 02 February 1993, Figs. 1-5.	18																		
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.																				
<table border="0"> <tr> <td>* Special categories of cited documents:</td> <td>*T</td> <td>later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</td> </tr> <tr> <td>*A* document defining the general state of the art which is not considered to be of particular relevance</td> <td>*X*</td> <td>document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</td> </tr> <tr> <td>*B* earlier document published on or after the international filing date</td> <td>*Y*</td> <td>document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</td> </tr> <tr> <td>*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</td> <td>*A*</td> <td>document member of the same patent family</td> </tr> <tr> <td>*O* document referring to an oral disclosure, use, exhibition or other means</td> <td></td> <td></td> </tr> <tr> <td>*P* document published prior to the international filing date but later than the priority date claimed</td> <td></td> <td></td> </tr> </table>			* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	*A* document defining the general state of the art which is not considered to be of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone	*B* earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art	*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A*	document member of the same patent family	*O* document referring to an oral disclosure, use, exhibition or other means			*P* document published prior to the international filing date but later than the priority date claimed		
* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention																		
A document defining the general state of the art which is not considered to be of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone																		
B earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art																		
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A*	document member of the same patent family																		
O document referring to an oral disclosure, use, exhibition or other means																				
P document published prior to the international filing date but later than the priority date claimed																				
Date of the actual completion of the international search 08 OCTOBER 1999		Date of mailing of the international search report 21 OCT 1999																		
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230		Authorized officer VY BUI <i>[Signature]</i> Telephone No. (703) 306-3420																		

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☐ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☐ **GRAY SCALE DOCUMENTS**
- ☐ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.